

PA	(INCYTE PHARM INC.
XX	
PI	Hillman JL, Corley NC, Patterson C;
XX	
DR	WPI: 2001-335067/35.
XX	
DR	P-PSDB; AAB85575.
XX	
PT	New substantially purified human glutathione peroxidase polypeptide,
PT	useful for diagnosing, treating or preventing reproductive disorders,
PT	immune disorders and cell proliferative or developmental disorders
XX	
PS	Example 1; Fig 1A-C; 26pp; English.
XX	
CC	This cDNA encodes a human glutathione peroxidase (Gpx6) polypeptide. The
CC	Gpx6 polypeptide is useful for diagnosing, treating or preventing
CC	disorders associated with expression of Gpx6, where the disorders are
CC	selected from reproductive disorders, immune disorders such as acquired
CC	immunodeficiency syndrome (AIDS), Addison's disease, adult respiratory
CC	distress syndrome, allergies, asthma, atherosclerosis, anemia, autoimmune
CC	hypothyroidism, bronchitis, diabetes mellitus, glomerulonephritis,
CC	Goodpasture's syndrome, gout, multiple sclerosis, myasthenia gravis,
CC	osteoporosis, rheumatoid arthritis, cancer, infections and trauma, and
CC	cell proliferative or developmental disorders such as arteriosclerosis,
CC	cirrhosis, psoriasis, cancer, Cushing's syndrome, and Sydenham's chorea.
CC	Gpx6 is also useful to produce antibodies, and to screen libraries of
CC	pharmaceutical agents to identify those which specifically binds Gpx6.
XX	
XX	Sequence 1072 BP; 275 A; 294 C; 270 G; 233 T; 0 other;

Query Match	100.0%;	Score 1072;	DB 22;	Length 1072;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches 1072; Conservative	0;	Mismatches	0;	Indels 0; Gaps 0;

QY	1	GAAGCGCCGACCTCCGGACAAACCCATGATGTCGCGGCGACGGTGGCAGCGCGTGGCTGCT	60
Db	1	gagcgccgcacactccgcgaacaagcatalggtgtgcgcgcagcttgccagtcgctgtcgtat	60
QY	61	CGTGTGGCGTGGCGCTGACGGCGACAGGACGACGACTTCCTACGACTTCGAAGCGGCTCAA	120
Db	61	ccctgttgagctcgtgcgcctgcgcgcagcagagcagacttctaagacttcaagcggctcaa	120
QY	121	CATCCGGGGGCAAACTGATGTGCTGGAGAAATACCGCGGATCGGTGTCCCTGGTGTGAA	180
Db	121	cattcgggggcaaacctgctgtcgtcgtgcgagaagtaccgcggatcgtgtcctgtgtgtga	180
QY	181	TGTGGCCAGCAGCTGCGGCTTCACAGACACACTACCAGACCTCGAGAGAGTGGAGGG	240
Db	181	tgtgcgcacagctgcgcgtcttcaacgacacactaccagacccctgcagagctgcagaag	240
QY	241	AGACCTGGGGCCCCGACCACTTCAGCTCGGCTCCGCCGCAACCAAGTTTGGCCAA	300
Db	241	agacctggggccccccacacacttcaagctgtcgtccttccctgcaccagttgtccaa	300
QY	301	GGAGCTTGACACCAACAGGAGATTGAGAGCTTTGCTGCCGCACTACAGTGTCTCAT	360
Db	301	ggagcttgacagcaacaagagatgtgagatgctgtcctgcgcacactacagtgctcatc	360
QY	361	CCCATGTTTACGAGATTGGAGTACCGGCTACTGTGTGTCCTCATCTGCTCAAGTACCT	420
Db	361	ccccatgtttaagaaagatgtgcagctaacgcgtactgtgtgccatccctgccttcaagta	420
QY	421	GGCCGACAGCTCGGGAAAGAGACCCACCTGGAAGCTTCGGAAGTACCTAGAGCCGAGA	480
Db	421	ggcccgacttcgggaagagagccacacgtgaacttcgtgaagtactctagttagcccgaga	480
QY	481	TGGAAAGTGTGAGGGCTTTGGACCAACTGTGTCACTGGAGGAGGTACAGACTCCAGAT	540
Db	481	tggaaagtggtgtagggcttgggaaccaactgtgtcagtgtagggaggtgcagatccagat	540
QY	541	CACAGCGCTCGTAGGAGACTCATCTTACTGAAGCGAAGAACTTTATACACCGGCTCT	600
Db	541	caacagcgctcgttgaagaacatcatcctactggaagcgaaagaaactataaacacacggtct	600

QY	601	CCCTCTCCACCACTCATCTCCCGCCACCTGTGTGGGGCTGACCAATGCAAACTCAATATG	660
Db	601	ccctctccacccactcatctccgcccacccgltgltggggctgtgaccaatgcaactcaaatg	660
QY	661	TGCTTCAAAGGAGAGACCCAGCTGCCTCCCTTATCTTTATGGCATTTGGTCCCAT	720
Db	661	tgcctcaaaaggagagccaccagcactccctcttcaactctatgcatgtgccat	720
QY	721	CATTCTGTGGGGAAAAATTTAGTATTTTGTATTTTGAATCTTTACAGCAACAATAG	780
Db	721	cattctgtggggaaaaatttctagtatttggattttgaactcttaagacaacaatag	780
QY	781	GACTCTCTGGGCATGAGACCTCTTACACAGTAATACACACGGATAGGACGCTTGC	840
Db	781	gaactctctgggcattgagacctcttgacccagtaaacacacagcgatatacgaaagcttgc	840
QY	841	CAACAAAATATTGTGGCAAAATAGAGTATTCAAGCAATATCTGCCACCAAGGCTTCT	900
Db	841	caacaaaatattgtggcaaaatagagtattcaagcaatattctgccaccagaagcttct	900
QY	901	GTAACCTGGGACCAATATTAATCTCATAGGGCTGTGTGAGATTAAGATGAANAACCTG	960
Db	901	gtaacctgggaccaatattatctcatagaggctgtgtgagattagatgaataacctg	960
QY	961	TGAAGTGCCCTAGGCAGTGCAGCAATAGAGCAATGAGGCAATCAATGAACAATTTTGGCATAT	1020
Db	961	tgaagtgccctaggcagtgccagcgaataatgaggagcatcattgtgcatat	1020
QY	1021	AAACCAAAAAATTAATCTGTTATCAATAAAACTTGCAATCAACATGAATTC	1072
Db	1021	aaacccaaaaataactcgtatcataataaaactgtatccacaatgaatttc	1072

RESULT 2

AA158027 standard; cDNA; 1100 BP.

AA
AC AAI58027;

DT 22-OCT-2001 (first entry)

AA	DE	Human polynucleotide SEQ ID NO 230.
AA	DE	Human polynucleotide SEQ ID NO 230.

Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer; KW

peripheral nervous system", central nervous system, etc.
KW Alzheimer's disease; Huntington's disease; haemostatic;

chemokinetic; thrombolytic; drug screening; arthritis; inflammation

leukaemia; SS:
KW
XX

OS Homo sapiens.
 YX

PN W0200153312-A1.

PD 26-JUL-2001

PF 26-DEC-2000; 2000WO-US34263.

PR 21-JAN-2000; 2000US-0488725.

PR 09-JUL-2000; 2000US-0598042.

PR 03-AUG-2000; 2000US-0653450.

PR 19-OCT-2000; 2000US-0693036.

XX	3	5	7	9	11	13	15	17	19	21	23	25	27	29	31	33	35	37	39	41	43	45	47	49	51	53	55	57	59	61	63	65	67	69	71	73	75	77	79	81	83	85	87	89	91	93	95	97	99
----	---	---	---	---	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----

(HIDE) INDEX INC
XX XX

PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;
PI Yang LL, Liu C, Huang Y, Chen M, Wu Z, Wang J, Zhang J;
PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;

P1 ZHANG YH, ZHANG F, GUO JICHEN N, DING WANGC XN/

1

XX WPI: 2001-442253/47.
DR P-PSDB: AAM38871.
XX
PT Novel nucleic acids and polypeptides, useful for treating disorders
PT such as central nervous system injuries -
PS Claim 1; SEQ ID NO 230; 10078bp; English.
XX
CC The invention relates to human nucleic acids (AA157798-AA161369) and
CC the encoded polypeptides (AAM38642-AAM42213) with nootropic,
CC immunosuppressant and cytostatic activity. The polynucleotides are useful
CC in gene therapy. A composition containing a polypeptide or polynucleotide
CC of the invention may be used to treat diseases of the peripheral nervous
CC system, such as peripheral nervous injuries, peripheral neuropathy and
CC localised neuropathies and central nervous system diseases, such as
CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
CC utilisation of the activities such as: immune system suppression,
CC Activin/Inhibin activity, chemotactic/chemokinetic activity, haemostatic
CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
CC assays for receptor activity, arthritis and inflammation, leukaemias and
CC C.N.S disorders.
CC Note: The sequence data for this patent did not form part of the printed
CC specification.
XX
SQ Sequence 1100 BP: 288 A; 305 C; 277 G; 230 T; 0 other:

Query Match 85.9%; Score 921; DB 22; Length 1100;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 1021; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GAGCGCGGACCTCGGAGCAAGCCATGTGGCGGAGCGTGGCGCGCGCTGCT 60
DB 23 gagcgccgcaaccccggaacaaagcaltgltggtgagcgtggtggtggtggtggt 82
QY 61 CCTGTGGGCTGGGCGCTGGCGCGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 120
DB 83 cctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 142
QY 121 CATCCGGGCGCAACTGTGTGCTGTGAGAGATACCGCGGATGCGTGTCCCTGGTGA 180
DB 143 catccgggcaaacactgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 202
QY 181 TGTGGCGACGAGTGTGGGCTTCAAGACAGACTACCGGCTGACAGAGAGAGAGAG 240
DB 203 tgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 262
QY 241 AGACCTGGGCGCGCGCGAGCTTCAACGTGCTGCTGCTGCTGCTGCTGCTGCTGCT 300
DB 263 agacctgggccccacacacttcaagtgctgtgtgtgtgtgtgtgtgtgtgtgtgt 322
QY 301 GAGCCTGACAGACAGAGAGAGATTTGAGAGCTTGTGCTGCTGCTGCTGCTGCT 360
DB 323 gtagcctgtgacagaaacagagaltgtagagcttgcgcgcgacacacacagtgctcat 382
QY 361 CCCGATGTTAGCAGATGTCAGTGCAGTGCAGTGCAGTGCAGTGCAGTGCAGTGCAG 420
DB 383 ccccatgttcaagaatgtgacgtacacgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 442
QY 421 GGGCCAGACTTGTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 480
DB 443 gggccagacttctgtggaagagagagagagagagagagagagagagagagagagag 502
QY 481 TGGAAAGGTGTGTGGGCTTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 540
DB 503 tggaaaggtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 562
QY 541 CACAGGCTGTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 600
DB 563 cacaggctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 622

QY 601 CCTCTTCACACACTTCATCCCGCCACCTGTGTGGGCTGTGACCAATGCAAACTCAATG 660
DB 623 cctcttcacacacttcattcccgccacactgtgtgtgtgtgtgtgtgtgtgtgtgtgt 682
QY 661 TGTCTTAAAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 720
DB 683 tgtcttaaaggagagagagagagagagagagagagagagagagagagagagagag 742
QY 721 CATCTGTGGGGGAAAAATTCATGATTTGATTTGATTTGATTTGATTTGATTTGAT 780
DB 743 catctctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 802
QY 781 GAACCTGTGGCGAGATGAGAGCTTGTGACAGAGTGAATCACCAGCGATAGCAAGCTT 840
DB 803 gaacctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 862
QY 841 CACAAAAAATGTGTGGCAATATGAAATATATCAAGCAATATATATATATATATAT 900
DB 863 caaaaaaatgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 922
QY 901 GTAACGTGGAGCAATGATTTACTCTAPAGGGCTGTGTGTGTGTGTGTGTGTGTGT 960
DB 923 gtaacctgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 982
QY 961 TGAAGTGTCTAGCAGAGTGTGACCAAGCAATAGAGGCAATCAATGAAATTTTGTGAT 1020
DB 983 tgaagtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 1042
QY 1021 AAA 1023
DB 1043 aaa 1045

RESULT 3
AA159813
ID AA159813 standard; cDNA; 1205 BP.
XX
XX AA159813;
AC
XX
DT 22-OCT-2001 (first entry)
XX
DE Human polynucleotide SEQ ID NO 3802.
XX
KW Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;
KW peripheral nervous system; neuropathy; central nervous system; CNS;
KW Alzheimer's; Parkinson's disease; Huntington's disease; hemostatic;
KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
KW chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
KW leukaemia; ss.
XX
OS Homo sapiens.
XX
PN WO200153312-A1.
XX
PD 26-JUL-2001.
XX
PF 26-DEC-2000; 2000MO-US34263.
XX
PR 21-JAN-2000; 2000US-0488725.
PR 25-APR-2000; 2000US-0552317.
PR 09-JUL-2000; 2000US-0598042.
PR 19-JUL-2000; 2000US-0620312.
PR 03-AUG-2000; 2000US-0653450.
PR 14-SEP-2000; 2000US-0662191.
PR 19-OCT-2000; 2000US-0693036.
PR 29-NOV-2000; 2000US-0727344.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;
PI Zhao QA, Zhou P, Goodrich R, Drmanac RT;

DR WPI: 2001-442253/47.
DR P-PSDB; AAM40657.
XX Novel nucleic acids and polypeptides, useful for treating disorders
PT such as central nervous system injuries -
XX

PS Claim 1; SEQ ID NO 3802; 10078bp; English.
XX
XX The invention relates to human nucleic acids (AA157798-AA161365) and
CC the encoded polypeptides (AAM38642-AAM42213) with nootropic,
CC immunosuppressant and cytostatic activity. The polynucleotides are useful
CC in gene therapy. A composition containing a polypeptide or polynucleotide
CC of the invention may be used to treat diseases of the peripheral nervous
CC system, such as peripheral nervous injuries, peripheral neuropathy and
CC localized neuropathies and central nervous system diseases, such as
CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
CC lateral sclerosis, and Shy-Drager syndrome. Other uses include the
CC utilization of the activities such as: Immune system suppression,
CC Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
CC assays for receptor activity, arthritis and inflammation, leukemias and
CC C.N.S disorders.
CC Note: The sequence data for this patent did not form part of the printed
CC specification.
CC
SQ Sequence 1205 BP; 324 A; 321 C; 289 G; 271 T; 0 other;

Query Match 85.9%; Score 921; DB 22; Length 1205;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 1021; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 1 GAGCGCGGCACCTCCGCGAACAAGCATGTGGCGCGAGCGGCGGCGTGGCTGCT 60
DB 2 gacgcgcgcaccccgcaacaagccatgctgcygcgcagcagcgtgcaagcgctgct 61
QY 61 CCGTGGCGCTCGCGCTCGCGCAGCAGAGCAGGACTTCTAGCATTCAGAGCGGCTCA 120
DB 62 cctgcygcgcgtgcygcgcgcagcagcagcagcagcagcagcagcagcagcagcagc 121
QY 121 CATCGCGGCGAAGCTGCTGCTGCTGAGAGTACGCGGATGCGGTGCTGCTGCTGTA 180
DB 122 catcgcgcgcacacgctgctgctgctgctgctgctgctgctgctgctgctgctgct 181
QY 181 TGTGCGCAGCAGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 240
DB 182 tctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgct 241
QY 241 AGACCTGGGCGCGCGCAGCTTACGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 300
DB 242 agacctggcgcccccacacttcaacgtctgctgctgctgctgctgctgctgctgct 301
QY 301 GGAGCGCTGACAGCAACAAGAGATTGAGAGCTTGTGCTGCTGCTGCTGCTGCTGCT 360
DB 302 ggaagcctgcaacaacaagagatgagagcttgcgcgcgcgcgcgcgcgcgcgcgcgc 361
QY 361 CCCCATGTTTACCAAGATTGACGACCGGTACTGGTGGCCATCTGCTTCAAGTACCT 420
DB 362 ccccatgtttagcaaatgtgacgtacacgctgctgctgctgctgctgctgctgctgct 421
QY 421 GCGCCAGACTTGTGGAGAGAGCCCACTGGAACCTTCTGGAAGTACTAGTACCCCA 480
DB 422 ggcgcagactcttggaagagagccacactggaactcttggaagtagtactagccccca 481
QY 481 TGGAAAGGTGTAGGGGCTTGGAGCCACTGTGCTGCTGCTGCTGCTGCTGCTGCTGCT 540
DB 482 tggaaaggtgtgtagggcttggagcccaactgtgtcagtggaggaagtagcagccagat 541
QY 541 CACAGCGCTGTGAGAGAGCTCATCTACTAGAGCAGAGAGATTAAACACCGCGTCT 600
DB 542 cacagcgctgtgtaggaagctcatctactagagcgagagacttaaccacacgctct 601
QY 601 CCTCTCCACCACTCATCCCGGCCACCTGTGTGGGGCTGACCATTGCAAACTCAATAG 660
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DB 602 cctctccacacactcatcccgccacactgtgtggtgagccaatgaaactcaatg 661
QY 661 TGCCTTCAAGGAGAGAGACCCACTGACTCTCTCTTACTCTTATGCGCATTTGGCCAT 720
DB 662 tgcctcaaaaggagagaccactgactctctctcttactcttattgcatgtgtccat 721
QY 721 CATCTCTGGGGGAAAAATCTAGTATTGATTATTTGATTTGATCTTACAGCAACAATAG 780
DB 722 catctgtggtgggaaaaatctgatttctgatttctgatttctgatttctgatttctgatt 781
QY 781 GAACCTCTGGCCAAATGAGAGCTTTGACCACTGAATCAACAGCCGATACGAAGCTTTGC 840
DB 782 gaactcccgccaatgagagctcttgaccagtgaaatccacgacgatacgagcgtctgc 841
QY 841 CACCAAAATGTGTGGCAATAGAGTATTCATCAACCATTAATCTCCCAACCAAGCTTCT 900
DB 842 caacaaaaatgtgtggcaaatagagtatatacaagaataatctcccaacaggtctct 901
QY 901 GTAACTGGAGCAATGATTAACCTCATAGGCGCTGTTGTGAGGATTAGATGAATACCTG 960
DB 902 gtaacttggaccatgatttactctatagggctgtgtgagagattagatgaataacctg 961
QY 961 TGAAGTGCCTTAGCGCAGTGCACGCCAATAGAGGCAATTCATGAACATTTTTCATAT 1020
DB 962 tgaagtgccttagcgagtgccagccaatagagagcatcattcaatctttgcatat 1021
QY 1021 AAA 1023
DB 1022 aaa 1024
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```
RESULT 4
AAC98225
ID AAC98225 standard; cDNA; 1321 BP.
XX
XX AAC98225;
AC
XX
XX 09-MAR-2001 (first entry)
DT
XX
XX Human colon cancer antigen nucleotide sequence SEQ ID NO:235.
DE
XX
XX Human; colon cancer; colon cancer antigen; diagnosis; detection;
KW Identification; cytostatic; cardioactive; neuroprotective; vulnery;
KW immunomodulatory; muscular; gynaecological; gastrointestinal;
KW nephrotropic; antiinfective; antibacterial; gene therapy; wound;
KW neural disorder; immune system disorder; muscular disorder;
KW reproductive disorder; gastrointestinal disorder; renal disorder;
KW infectious disease; cardiovascular disorder; ss.
XX
XX Homo sapiens.
OS
XX
XX WO20005351-A1.
PN
XX
XX 21-SEP-2000.
PD
XX
XX 08-MAR-2000; 2000WO-US05883.
PF
XX
XX 12-MAR-1999; 99US-0124270.
PR
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PA
XX
XX Rosen CA, Ruben SM;
PI
XX
XX WPI: 2000-587534/55.
DR
XX
XX P-PSDB; AAB53468.
PT
XX
XX Colon cancer associated gene sequences, referred to as colon cancer
PT antigens, useful for the treatment, prevention, and diagnosis of colon
PT disorders such as colon cancer -
PS
XX
XX Claim 1; Page 656-657; 2104pp; English.
```

CC AAC97991 to AAC98763 encode the human colon cancer associated proteins,
 CC called human colon cancer antigens, given in AAB5324 to AAB54006. The
 CC human colon cancer antigens can have cytosolic, cardiovacular, muscular,
 CC neuroprotective, immunomodulatory, gynaecological, gastrointestinal,
 CC and vulnerrary, nephrotropic, anti-infective and antibacterial activities, and
 CC can be used in gene therapy. The colon cancer antigen polynucleotides,
 CC proteins and antibodies of the proteins are useful for the prevention,
 CC treatment and diagnosis of colon disorders, such as colon cancer. The
 CC polynucleotides may be used in diagnostics and research, such as for
 CC chromosome identification, and as hybridisation probes. The proteins
 CC may also be used to prevent diseases such as neural disorders, immune
 CC system disorders, muscular disorders, reproductive disorders,
 CC gastrointestinal disorders, wounds, renal disorders, infectious
 CC diseases, and cardiovascular disorders. AAC98764 to AAC98772 and
 CC AAB54007 represent sequences used in the exemplification of the present
 CC invention.

XX Sequence 1321 BP; 420 A; 326 C; 296 G; 276 T; 3 other;

Query Match 85.9%; Score 921; DB 21; Length 1321;
 Best Local Similarity 99.8%; Pred. No. 0;
 Matches 1021; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GAGCGCCGACCTCCGGAACAGCCATGCTGGCGGACGCTGGACGCGCTGGCTGCT 60
 DB 5 gacgcgcacactccgcgaacaaagcatgctgctgagcagcgcgcgcgcgcgcgcgcgc 64
 QY 61 CCTGTGGGTGGGGCTGGCGGACGAGAGAGACTTCTACGACTTAAAGCGGTCAA 120
 DB 65 cctgtggtgctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgct 124
 QY 121 CATCCGGGGCAACATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 180
 DB 125 catccggggcaaacatgctgctgctgctgctgctgctgctgctgctgctgctgctgct 184
 QY 181 TGTGGCCACGAGTGGCGCTTACAGACGACGACGACGACGACGACGACGACGACG 240
 DB 185 tgtggccacgagtggcgcttaccagacgacgacgacgacgacgacgacgacgacgacg 244
 QY 241 AAGACCTGGGGCCGACCTTCAAGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 300
 DB 245 aagacctggggccgacaccttcaacgctgctgctgctgctgctgctgctgctgctgctg 304
 QY 301 GGAGCCTGACGACCAACAGAGATTGAGAGCTTGGCTCCGACGCTTACAGTGTCTCAT 360
 DB 305 ggaagcctgacgacaaacagagattgagagcttggccgcgcgcgcgcgcgcgcgcgcgc 364
 QY 361 CCCCATGTTTACAGATGTCAGTACCGGTACTGGTGGCCATCTGCTTCAAGTACCT 420
 DB 365 ccccatgtttacgaattgtcagtcacgcgtactgtgcccattcctcctcaagtaacct 424
 QY 421 GGCCGAGACTTCTGGGAAGAGCCGACCTGGAATCTCTGGAGTACTAGTACCCCA 480
 DB 425 ggccgagacttctgggaagagccgacccctggaactctggaagtagtactgagccca 484
 QY 481 TGGAAAGTGTAGGAGCTTGGGACCAACTGTCTAGTGAAGAGAGTGAAGTCCAGAT 540
 DB 485 tggaaagtgtagaggcttgggacccaactgtctagtgagggagagtagtaagaccagat 544
 QY 541 CACAGCGCTGTGAGAGACTCTACTGAGCGAGAGAGACTTAAACACCGGCTCT 600
 DB 545 cacagcgctgtgagagactctactgagcgagagacttaaacacacgcgctct 604
 QY 601 CCTCCGACGACGCTGATCCGCGCCGCTGTGGGGGCTGACCAATGCAACGCAATAG 660
 DB 605 cctccgacgacgctgattccgcgcacccctgtggtgggcgacaaagcaaaccaatag 664
 QY 661 TGGCTTAAAGGAGAGACCCACTGACTCTCTCTTACTCTTAAAGCAATTGGTCCAT 720
 DB 665 tggctttaaaggagagaccactgactctctccttactctactgacatgtgtccat 724
 QY 721 CATTTCTTGTGGGGAAAAATTTCTAGTATTTTGTATTGTAATCTTTACAGCAACAATAG 780

DB 725 catctctggtgggaaataatctagatatttgattatgtaacttaccagcaacaatag 784
 QY 781 GAACCTCTGGCCAAATGAGAGCTTTGACAGTCAATACACAGCCGCTTACGAGCTTTCG 840
 DB 785 gaactctgtgcaaatgagagctcttgaccagtgtaaccacagcgataagaaagcttgc 844
 QY 841 CACCAAAATGTTGTGCAATATAGATATATCAAGCAATATCTCCACCAAGGCTTCT 900
 DB 845 caacaaaatgtgtgcaaatagaaatagataatacaagcaataatctccacccaagcttct 904
 QY 901 GTAACCTGGGACCAATGATTACCTCATAGGCTGTTGTGAGATTAGATGAATACCTG 960
 DB 905 gtaactgggaccaaataatgattacctaataagagctgtgtgagattagatgaataacc 964
 QY 961 TGAAGTGGCTAGCAGTGGCCAGCCAAATAGAGGCAATTCATGACATTTTTCATAT 1020
 DB 965 tgaagtggctagcagtgccagccaataagagagcattcaatgaacatttttgatcat 1024
 QY 1021 AAA 1023
 DB 1025 aaa 1027

RESULT 5
 AAH72778
 ID AAH72778 standard; cDNA; 1511 BP.
 XX
 AC AAH72778;
 XX
 DT 19-SEP-2001 (first entry)
 XX
 DE Human cervical cancer marker nucleic acid 4052.
 XX
 KW Cervical cancer; cytostatic; pre-malignant condition; gene therapy; ss.
 XX
 XX Homo sapiens.
 PN WO200142467-A2.
 XX
 PD 14-JUN-2001.
 XX
 PF 08-DEC-2000; 2000WO-US3312.
 XX
 PR 08-DEC-1999; 99US-0169681.
 PR 21-DEC-1999; 99US-0171350.
 PR 14-MAR-2000; 2000US-0189315.
 PR 12-MAY-2000; 2000US-0203791.
 PR 09-JUN-2000; 2000US-0210600.
 PR 21-JUL-2000; 2000US-0220114.
 XX
 PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
 XX
 PI Schlegel R, Deeds J, Berger A, Zhao X;
 DR
 WIPI: 2001-375006/39.
 PT New isolated nucleic acid for diagnosing and treating cervical cancer
 CC and for assessing and detecting compounds for treating the cancer -
 XX
 PS Claim 1; Page 845-847; 1051pp; English.
 XX
 CC The invention relates to novel genes (AAH68727-AAH73383) associated with
 CC cervical cancer with cytostatic activity. The nucleic acids and encoded
 CC polypeptides are useful: to assess if a patient is afflicted with
 CC cervical cancer or has a pre-malignant condition; to monitor the
 CC progression of cervical cancer or a premalignant condition in a patient;
 CC and to select and/or assess the efficacy of a compound or therapy for
 CC inhibiting cervical cancer in a patient. The nucleic acids may also be
 CC useful for gene therapy.
 XX
 SQ Sequence 1511 BP; 392 A; 421 C; 378 G; 314 T; 6 other;

Matches 1069; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

```
OY 1 GAGCGCGCCACCTCCGGAACAAGCCATGATGGCGGAGCGGAGCGGCGCTGCT 60
    |||||||
DB 15 gacgcgcgcacctcccgaaacaagcatggtgcygcgaagcggcgagcgtgtgctgt 74
OY 61 CCTGTGGGCTGGCGCTGGCGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 120
    |||||||
DB 75 cctgtggtgctggtgctggtgctggtgctggtgctggtgctggtgctggtgct 134
OY 121 CATCCGGGGCAACATGCTGCTGGAGAAATACCGCGATGGGTGCTGCTGCTGAA 180
    |||||||
DB 135 catccggggcaaacctgtgtcgtggaagatccggtgctgctcctgtgtgtgaa 194
OY 181 TGTGGCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 240
    |||||||
DB 195 tgtggtcagcaggtgtggtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgtgt 254
OY 241 AGACCTGGGGCCCCACACACTTCAACGTGCTGCTGCTGCTGCTGCTGCTGCT 300
    |||||||
DB 255 agacctgggccccaccaccttaacgtgtcgtcctccctgcaaccagttgtgccaac 314
OY 301 GGAGCCTGACACACAACAAGAGATGACAGCTTGGCTCCGACCTACAGTGTCTATT 360
    |||||||
DB 315 gtagcctgacagcaacaagagattgagagcttggccgcgcacactacagttctcat 374
OY 361 CCCCATGTTTACAGATGTCAGTCAACGGTACTGCTGCTGCTGCTGCTGCTGCT 420
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DB 375 ccccatgttttagcaagattgacagtcacggttactgtgtccatccctgcttaagta 434
OY 421 GGCCCAAGACTTCTGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 480
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DB 435 ggcacagactctgtggaagagccacactgtgaactctgtgaagtagcctcagc 494
OY 481 TGGAAAGGTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 540
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DB 495 tggaaaggtgtagagagctgtggaacacacgtgtcagtggaagagtlacagacc 554
OY 541 CACAGCGCTGTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 600
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DB 555 cacagcgctgtgaggaagctacactcctactgaagcgaagactttaaccacgcgt 614
OY 601 CCTCCTCCACACACTTATCCCGCCACCTGTGTGGGCTGACCAATGCAACTCAAT 660
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DB 615 cctcctccacacactatccgcacctgtgtgtggtggtggtggtggtggtggtg 674
OY 661 TGGTTCAAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 720
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DB 675 tgcctcaaaaggagagaccacacgtactcctccttactcttactgtgcaatgtgt 734
OY 721 CATTTCTTGGGGGAAAAATTTAGTATTGATTGATTGATTGATTGATTGATTGATT 780
    |||||||
DB 735 catctctgtgggggaaaaattctagatattgtatttgaatcttaacagcaacaat 794
OY 781 GAACCTCTGGCCCAATGAGAGCTTTGACCAAGTAATCACCAGCCGATAGACGT 840
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DB 795 gaactcctgtgccaatgagagctcttgacacagtlacaccagcgaacagtcgtgc 854
OY 841 CAACAAAAATGTTGCAATATAGATATATCAAGCAATATATCCACCCCAAGGCTCT 900
    |||||||
DB 855 caacaaaaatgtgtgcaaatatagataatacaagaataatctccacccaaggctct 914
OY 901 GTAACCTGGAGCAATGATTACCTCATAGAGGCTGTTGTGAGATTAGATGAATAC 960
    |||||||
DB 915 gtaacctggagcaaatgattactcctagagctgtgtgtgtgtgtgtgtgtgtgt 974
OY 961 TGAATGCTGCTAGGAGTGTGCGCAAGCAATAGAGGCAATCAATGAACATTTTGCAT 1020
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DB 975 tgaatgtcctagcagtgccagccaatagagatcaatgaacatcttltgtcatat 1034
OY 1021 AAACCAAAAAATGTTGTTATCAATAAAACTTGATCCCAATGAAATTTTC 1072
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DB 1035 aaaccaaaaaataactgttatacaataaaaaactgtcatccaacatgaatttc 1086
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RESULT 7
AAZ65013
ID AAZ65013 standard; cDNA; 1227 BP.
XX
AC AAZ65013;
XX
DT 05-APR-2000 (first entry)
XX
DE Membrane-bound protein PRO828 encoding cDNA.
XX
KW Membrane-bound polypeptide; PRO polypeptide; LDL receptor; TIE ligand;
pharmaceutical; receptor immunoadhesin; gene mapping; ss.
OS Homo sapiens.
XX
PN WO9963088-A2.
XX
PD 09-DEC-1999.
XX
PF 02-JUN-1999; 99MO-US12252.
XX
PR 02-JUN-1998; 98US-0087607.
PR 02-JUN-1998; 98US-0087609.
PR 02-JUN-1998; 98US-0087759.
PR 03-JUN-1998; 98US-0087827.
PR 04-JUN-1998; 98US-0088021.
PR 04-JUN-1998; 98US-0088025.
PR 04-JUN-1998; 98US-0088028.
PR 04-JUN-1998; 98US-0088029.
PR 04-JUN-1998; 98US-0088030.
PR 04-JUN-1998; 98US-0088033.
PR 04-JUN-1998; 98US-0088326.
PR 05-JUN-1998; 98US-0088167.
PR 05-JUN-1998; 98US-0088202.
PR 05-JUN-1998; 98US-0088212.
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PR 10-JUN-1998; 98US-0088722.
PR 10-JUN-1998; 98US-0088730.
PR 10-JUN-1998; 98US-0088734.
PR 10-JUN-1998; 98US-0088738.
PR 10-JUN-1998; 98US-0088740.
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PR 10-JUN-1998; 98US-0088742.
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PR 10-JUN-1998; 98US-0088825.
PR 10-JUN-1998; 98US-0088826.
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PR 11-JUN-1998; 98US-0088876.
PR 12-JUN-1998; 98US-0089090.
PR 12-JUN-1998; 98US-0089105.
PR 16-JUN-1998; 98US-0089440.
PR 16-JUN-1998; 98US-0089512.
PR 16-JUN-1998; 98US-0089514.
PR 17-JUN-1998; 98US-0089532.
PR 17-JUN-1998; 98US-0089538.
PR 17-JUN-1998; 98US-0089598.
PR 17-JUN-1998; 98US-0089599.
PR 17-JUN-1998; 98US-0089600.
PR 17-JUN-1998; 98US-0089653.
PR 18-JUN-1998; 98US-0089801.
PR 18-JUN-1998; 98US-0089907.
PR 18-JUN-1998; 98US-0089908.
PR 19-JUN-1998; 98US-0089947.
PR 19-JUN-1998; 98US-0089948.
PR 19-JUN-1998; 98US-0089952.
PR 22-JUN-1998; 98US-0090246.
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PR 22-JUN-1998; 98US-0090252;
PR 22-JUN-1998; 98US-0090253;
PR 23-JUN-1998; 98US-0090349;
PR 23-JUN-1998; 98US-0090355;
PR 24-JUN-1998; 98US-0090421;
PR 24-JUN-1998; 98US-0090431;
PR 24-JUN-1998; 98US-0090435;
PR 24-JUN-1998; 98US-0090444;
PR 24-JUN-1998; 98US-0090445;
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PR 24-JUN-1998; 98US-0090472;
PR 24-JUN-1998; 98US-0090535;
PR 24-JUN-1998; 98US-0090538;
PR 24-JUN-1998; 98US-0090540;
PR 24-JUN-1998; 98US-0090557;
PR 25-JUN-1998; 98US-0090676;
PR 25-JUN-1998; 98US-0090678;
PR 25-JUN-1998; 98US-0090688;
PR 25-JUN-1998; 98US-0090690;
PR 25-JUN-1998; 98US-0090691;
PR 25-JUN-1998; 98US-0090694;
PR 25-JUN-1998; 98US-0090695;
PR 25-JUN-1998; 98US-0090696;
PR 26-JUN-1998; 98US-0090862;
PR 26-JUN-1998; 98US-0090863;
PR 01-JUL-1998; 98US-0091358;
PR 01-JUL-1998; 98US-0091360;
PR 01-JUL-1998; 98US-0091544;
PR 02-JUL-1998; 98US-0091478;
PR 02-JUL-1998; 98US-0091486;
PR 02-JUL-1998; 98US-0091519;
PR 02-JUL-1998; 98US-0091626;
PR 02-JUL-1998; 98US-0091628;
PR 02-JUL-1998; 98US-0091633;
PR 02-JUL-1998; 98US-0091646;
PR 02-JUL-1998; 98US-0091673;
PR 07-JUL-1998; 98US-0091978;
PR 07-JUL-1998; 98US-0091982;
PR 09-JUL-1998; 98US-0092182;
PR 10-JUL-1998; 98US-0092472;
PR 20-JUL-1998; 98US-0093339;
PR 30-JUL-1998; 98US-0094651;
PR 04-AUG-1998; 98US-0095282;
PR 04-AUG-1998; 98US-0095285;
PR 04-AUG-1998; 98US-0095301;
PR 04-AUG-1998; 98US-0095302;
PR 04-AUG-1998; 98US-0095318;
PR 04-AUG-1998; 98US-0095321;
PR 04-AUG-1998; 98US-0095325;
PR 10-AUG-1998; 98US-0095916;
PR 10-AUG-1998; 98US-0095929;
PR 11-AUG-1998; 98US-0096012;
PR 11-AUG-1998; 98US-0096146;
PR 12-AUG-1998; 98US-0096329;
PR 17-AUG-1998; 98US-0096757;
PR 17-AUG-1998; 98US-0096766;
PR 17-AUG-1998; 98US-0096768;
PR 17-AUG-1998; 98US-0096773;
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PR 17-AUG-1998; 98US-0096894;
PR 17-AUG-1998; 98US-0096895;
PR 17-AUG-1998; 98US-0096897;
PR 18-AUG-1998; 98US-0096949;
PR 18-AUG-1998; 98US-0096950;
PR 18-AUG-1998; 98US-0096959;
PR 18-AUG-1998; 98US-0096960;
PR 18-AUG-1998; 98US-0097022;
PR 19-AUG-1998; 98US-0097141;
PR 20-AUG-1998; 98US-0097218;
PR 24-AUG-1998; 98US-0097661;

PR 26-AUG-1998; 98US-0097951;
PR 26-AUG-1998; 98US-0097952;
PR 26-AUG-1998; 98US-0097954;
PR 26-AUG-1998; 98US-0097955;
PR 26-AUG-1998; 98US-0097971;
PR 26-AUG-1998; 98US-0097974;
PR 26-AUG-1998; 98US-0097978;
PR 26-AUG-1998; 98US-0097979;
PR 26-AUG-1998; 98US-0097986;
PR 31-AUG-1998; 98US-0098014;
PR 31-AUG-1998; 98US-0098525;
PR 16-SEP-1998; 98US-0100634;
PR 12-JAN-1999; 99US-0115565;
XX
XX (GETH) GENENTECH INC.
XX
XX Baker K, Chen J, Goddard A, Gurney AL, Smith V, Watanabe CK;
PI Wood WI, Yuan J;
XX
XX WPI: 2000-072883/06.
DR P-PSDB: AAY66677.
XX
XX Membrane-bound proteins and related nucleotide sequences -
PS
PS Claim 2; Fig 119; 822pp; English.
XX
XX
XX The invention provides membrane-bound PRO polypeptides and
CC polynucleotides encoding them. The PRO sequences of the invention were
CC identified based on extracellular domain homology screening. The PRO
CC sequences have homology with proteins including LDL receptors, the
CC ligands and various enzymes. The membrane-bound proteins and receptor
CC molecules are useful as pharmaceutical and diagnostic agents. Receptor
CC immunoadhesins, for instance, can be used as therapeutic agents to block
CC receptor-ligand interactions. The membrane-bound proteins can also be
CC employed for screening of potential peptide or small molecule inhibitors
CC of the relevant receptor/ligand interaction. The PRO encoding sequences
CC are useful as hybridization probes, in chromosome and gene mapping and in
CC the generation of antisense RNA and DNA. PRO nucleic acid sequences
CC will also be useful for the preparation of PRO polypeptides, especially
CC by recombinant techniques.
XX
XX Sequence 1227 BP; 331 A; 325 C; 293 G; 278 T; 0 other;
SQ
Query Match 85.4%; Score 916; DB 21; Length 1227;
Best Local Similarity 99.7%; Pred. No. 0;
Matches 1066; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 4 GCCGCCACCTCCGGAACAGCCATGTGGCGGAGGAGGCGGCGGCTGCTCC 63
DB 12 gccgcacccctccggaacagccatgttgcgcgagcggtgagcggtgctgcctc 71
QY 64 GTGGGCTGGGCTGCGCGCAGCAGCAGAGACTTCTACGACTTCAAGCGGTCAACAT 123
DB 72 gtggctggtggctgctgcgcagcagagagcaggtcttcaagcttaagcggtcaacat 131
QY 124 CCGGGGCAAACTGCTGCTGCGTGAAGAATACCGCGATGCTGCTGCTGCTGATGT 183
DB 132 ccggggcaaaactgtgtcgtcgtgagaagtacgcgcgagatcggttccctggtgtaagt 191
QY 184 GGGCAGCAGTGGGCTTACAGACACGACTACCGAGCCCTGACAGAGTGGAGAGA 243
DB 192 ggcacagcagtgctgttccacacacagcactacagccctgcagagctgagcgaga 251
QY 244 CCTGGGCCCCCACCACCTTCAACGTGCTGCTGCCCTTCCCTGCAACAGTTTGGCAACAGA 303
DB 252 cctgggccccaccacttcaacgtgtcgtccttcccttgcacacagtttggccaacaga 311
QY 304 GCCTGACAGCAACAGAGAGATTGAGAGCTTGGCTGCGCGCACTACAGTGTCTCATTTCC 363
DB 312 gcttgaacagcaacagagattgagagcttgcgcgcgcacactaagttgtcattccc 371
QY 364 CATGTTAGCAAGATTGCATCAACCGGTACTGTTGCCCATCTGCTTCAAGTACTGGC 423

Query Match 85.4%; Score 916; DB 22; Length 1227;
 Best Local Similarity 99.7%; Pred. No. 0;
 Matches 1066; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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OY 4 GCGGCCACCTCCGGAACAAGCCATGCTGCGCGACGCTGGCAGCGCGCTGCTCCT 63
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DB 12 gccgcacacctccgacaagaacccaatggtgctgacgagtgacgagcgtgctgctcc 71
    |||||
OY 64 GTCGGCTGCGGCTGGCGGACGAGGAGGAGCTTCTAGACTTCAAGCGGTCACAT 123
    |||||
DB 72 gtgggctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgct 131
    |||||
OY 124 CCGGGCCAAACTGCTGCTGCTGAGAGTACCGCGATGCTGCTGCTGCTGCTGCT 183
    |||||
DB 132 ccggggcaaaactgctgctgctgctgctgctgctgctgctgctgctgctgctgct 191
    |||||
OY 184 GCGCCAGCAGTGGGCTTTCACAGACCCAGCTACCGCCCTGACAGAGCTGCGCGAGA 243
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DB 192 ggcacagcagatgctgctgctgctgctgctgctgctgctgctgctgctgctgct 251
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OY 244 CCGGGGCCCCACACTTCAAGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 303
    |||||
DB 252 cctgggccccccacacttaacgtgctgctgctgctgctgctgctgctgctgctgct 311
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OY 304 GCGTCAGCAGCAACAGAGATTTAGACCTTTGCTGCGCCAGCTTACAGTCTTATCCC 363
    |||||
DB 312 gccgcagacaagaagagatgagagcttgcgcgcgcgcacacagtgctcatccc 371
    |||||
OY 364 CAGTTTACGAGATGTCAGTGCAGTGCAGTGCAGTGCAGTGCAGTGCAGTGCAGTGC 423
    |||||
DB 372 cagtgtagcaagatgctgctgctgctgctgctgctgctgctgctgctgctgctgct 431
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OY 424 CCAGACTTCTGGAAGAGACCCACTTGGAACTTCTGGAAGTACTAGTACGCCAGATGG 483
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DB 432 ccagactcttggaagagagccacttgaactcttggaagtagcagcccaagag 491
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OY 484 AAAGTGTAGGGCTTGGAGCCACACTGTCTCAAGTGGAGAGGTCAGACATCCAGATCAC 543
    |||||
DB 492 aaagtgtagagagcttggagcccaactgtgctgtagagagtgcaaccaccatcac 551
    |||||
OY 544 AGCGCTGTGAGGAGTCTCATCTTACTGAGAGAGAGTCTTATACCAACCGCTCTCT 603
    |||||
DB 552 agcgctgtgagagagctcactcactcactcactcactcactcactcactcactcact 611
    |||||
OY 604 CCTCCACCACTCACTCCCGCCCACTGCTGGGCTGACCAATGCAATCAATGCTGTC 663
    |||||
DB 612 cctccacacactcactcctgctgctgctgctgctgctgctgctgctgctgctgctgct 671
    |||||
OY 664 TTCAAGGAGAGACCACTGACTCTCTCTCTTACTTATGCTTATGCTTATGCTTATG 723
    |||||
DB 672 ttcaaaaggagagacacactcactcactcactcactcactcactcactcactcact 731
    |||||
OY 724 TCTGTGGGGGAAAAATTTCTAGTATTTGATTTGATTTGATTTGATTTGATTTGATTT 783
    |||||
DB 722 tctgtgggggaaaaaattctagatatttgatatttgatatttgatatttgatatttgat 791
    |||||
OY 784 CTCCTGGCAATGAGAGCTTGTGACAGGATCACACAGCCAGTACGAGCTTGTGCCAA 843
    |||||
DB 792 ctcctggccaagagagctctgacacagtgatcacccagcgatagaaagcgtctgcca 851
    |||||
OY 844 CAAAAATGTGTGGCAATAGAGTATATCAACCAATATCTCCACCCAGGCTTGTGTA 903
    |||||
DB 852 caaaaatgtgtggcaaatagagatatacaagaataatccaccacgaagctcttgta 911
    |||||
OY 904 AACTGGGACCAATGATTTACCTATAGGCTGTTGTGAGATTAAGTGAATACCGTGA 963
    |||||
DB 912 aactgggaccaaagtattactcattagagctgtgtgagatagatagaaataacccgtg 971
    |||||
OY 964 AAGTGCTTGGAGTCCAGCCAAATAGGAGCATTCATTTGATTTGATTTGATTTGATTT 1023
    |||||
DB 972 aagtgcttggagctgctgctgctgctgctgctgctgctgctgctgctgctgctgctgct 1031
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OY 1024 CCAAAAAATTAACCTGTTATCAATAAAAACTTGATCCAAATGATTTTC 1072
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DB 1032 ccaaaaataactgttatacaataaaacttgcatccacaatgaatttc 1080
 |||||

RESULT 9

AA#44159
 ID AA#44159 standard; cDNA; 1227 BP.

AA#44159;

02-APR-2001 (first entry)

Human PRO828 (UNQ469) nucleotide sequence SEQ ID NO:188.

Human; secreted and transmembrane protein; PRO: cytosolic;
 cell death; cancer; chromosomal mapping; gene mapping; tissue typing;
 diagnostic assay; ss.

Homo sapiens.

WO200073454-A1.

07-DEC-2000.

30-MAR-2000; 2000WO-US08439.

02-JUN-1999; 99WO-US12252.
 23-JUN-1999; 99US-0141037.
 07-JUL-1999; 99US-0143048.
 20-JUL-1999; 99US-0144758.
 26-JUL-1999; 99US-0145698.
 28-JUL-1999; 99US-0146222.
 17-AUG-1999; 99US-0149396.
 15-SEP-1999; 99WO-US21090.
 15-SEP-1999; 99WO-US21547.
 08-OCT-1999; 99US-0158663.
 30-NOV-1999; 99WO-US28313.
 01-DEC-1999; 99WO-US28301.
 16-DEC-1999; 99WO-US30095.
 20-DEC-1999; 99WO-US30911.
 05-JAN-2000; 2000WO-US00219.
 06-JAN-2000; 2000WO-US00376.
 11-FEB-2000; 2000WO-US03565.
 18-FEB-2000; 2000WO-US04341.
 22-FEB-2000; 2000WO-US04414.
 24-FEB-2000; 2000WO-US04914.
 24-FEB-2000; 2000WO-US05004.
 02-MAR-2000; 2000WO-US05841.
 15-MAR-2000; 2000WO-US06884.
 20-MAR-2000; 2000WO-US07377.

(GETH) GENENTECH INC.

Ashkenazi AJ, Baker KP, Botstein D, Desnoyers L, Eaton DL,
 Ferrara N, Fong S, Gerber H, Gertlsen ME, Goddard A, Godowski PJ,
 Grimaldi CJ, Gunney AL, Kijavini RJ, Napier MA, Pan J, Paoni NF,
 Roy MA, Stewart TA, Tumas D, Watanabe CK, Williams PM, Wood WT,
 Zhang Z;

WPI: 2001-032160/04.

P-PSDB: AMB65200.

PRO polynucleotides used to produce polypeptides used to target
 bioactive molecules such as toxins, radiolabels or antibodies, to
 specific cells, to cause targeted cell death -

Claim 2: Fig 119; 935pp; English.

The present invention describes human secreted and transmembrane PRO
 proteins. The PRO proteins have cytosolic activity. The PRO proteins
 can be used for targeted delivery of bioactive molecules, such as
 toxins, radiolabels or antibodies, that cause cell death. PRO nucleotide
 sequences, and their fragments, can be used as hybridisation probes. In

CC chromosomal and gene mapping, and in the generation of anti-sense RNA
CC and DNA. They may also be used to produce transgenic animals which are
CC used to develop and screen therapeutically useful reagents. The PRO
CC nucleotide and protein sequence can be used for tissue typing and in
CC treating cancer. Anti-PRO antibodies can be used in diagnostic assays.
CC AA644220 to AA644470 represent PCR primers and hybridisation probes used
CC in the isolation of human PRO sequences. AA644087 to AA644269 and
CC AA665154 to AA665300 represent human PRO polynucleotide and protein
CC sequences given in the exemplification of the present invention.

SQ Sequence 1227 BP; 331 A; 325 C; 293 G; 278 T; 0 other;

Query Match	85.4%	Score 916	DB 22	Length 1227
Best Local Similarly	99.7%	Pred. No. 0		
Matches 1066	Conservative	0	Mismatches 3	Indels 0
				Gaps 0

OY	4	GCCTCCACCTCTCGGAAACAAGCAGCATGCTGGCGGGGCGACGCTGGCAGCGCGCTGCTCCT	63
Db	12	gcgcgcacctctccggaacaagaagcaltggtgycgycgaacggtgycagcgagctggtgtctcct	71
OY	64	GTGGGCTCCGGCTCGCGCGACGACGAGACAGATTCTTACGACTTCAAGCGGCTCAACAT	123
Db	72	gtgggctctgcygcccgcgcgcgcagcagcgagcagacttctcaagtcgaagcggltcaaat	131
OY	124	CCGGGGGCAAACTGGTGTGCTGCGTGGAGAAGTACCGGGATCGGTGTCCCTGGTGTGATGT	183
Db	132	ccggggcaaaactggtgtctgctgtagaagtaaccggagatcggtgtcctgtgtgtgtagt	191
OY	184	GGCCACGACAGTGGCGGCTTACACAGCCAGCACTACCGAGCCCTGTACGACGCTGCAGGAGA	243
Db	192	ggccagcgagtgcygcttcaacaagccagcctaacccgagcccttcagagagcttgcaagcgaga	251
OY	244	CGTGGGCCCCCACCACCTTTCACGTCGCTGCTCCCTCGCAACACAGTTTGCCACACAGA	303
Db	252	ccgtggccccccacactttaacgctgcgcctctcccttcgaacacagtttgccaacaagga	311
OY	304	GCCTGCACAGCAACAAGAGATTGAGACCTTGGCTGCCGACCTACAGTGTCTCATPCC	363
Db	312	gcctgcacagcaacaagagatltgagactltgccccgcagcactaacaglttccattccc	371
OY	364	CATGTTTGGCAAGATTGAGACACCGGTTACGGGTGGCCATCTCGTCTTCAAGTAACTCGCG	423
Db	372	catgtttgcaagatttgaaagcaccggtuacggtggcccactccgacctcaagtaactcggc	431
OY	424	CCAGACTCTTGGGAAGAGGCCACCTCTGGAATTCTCGAAGTACTAGTACCCACAGATGG	483
Db	432	ccagactctctggagaagagccacactctggaactctctggaagtaactagtagccccagatg	491
OY	484	AAAGTGGTAAAGGGCTTGGGACCACACTGTGTCAGTGGAGAGGTCAAGACTCCAGATCAC	543
Db	492	aaagtggtlaagggctctgtagcccaactgtgtcagltgagagagtgacagcccaagatcac	551
OY	544	AGCGTCTGTAGGAAGCTCATCTACACTGAAGCGAGAGACTTTTAAACACCGGTCMCC	603
Db	552	agcgctctgtaggaagctcatctccactcgaagcaggaagacttatcaacccgcgctcct	611
OY	604	CCTCCACCACTCATCCCGGCCACCTGTGTGGGGCTTACCAATGCAATCAATGGTGC	663
Db	612	ctctccacactcatctccgcgccacactgtgtggygcgtgaccaatgcaactcaaatgtgc	671
OY	664	TTTCAAGGGAAGAACCCACTGACTCTCTCTCTTACTCTTAAGCCATTGGTCCCATCAT	723
Db	672	ttcaaaaggagaagaccacatgactctctctcttactcttctatgcaatgtgtccatcat	731
OY	724	TCTTGTGGGGAAAAATCTAGTATTTTGTGATTATTCAATCTTAACACCAATATGGA	783
Db	732	tctgtggyggaaaaaattctcgtattttgttatatttgaactctcaagcaacaatatgga	791
OY	784	CTCTCGGCATATGAGAGCTTTGACCAAGTGATATCCAGCCGATAGCAAGCTTTGGCAA	843
Db	792	ctctcgcgcaaatgtgagctcttgccagtgtaataccaagccgataacgaacgctctccaa	851

OY	844	CAAAAATGTGTGGCAAAATAGAAAGTATTTTCACGAATATATTTCCACCAAGGCTTCTGTA	903
Db	852	caaaaatgtgtggaacaatagaaatatacaagaacaataatctccaccccaagctctctgta	911
OY	904	AACGTGGACCAATATATTTACCTTCATAGGGCTGTGTAGGATTTAGATGAATATACCTGTGA	963
Db	912	aaacttgagccaaatatttaacctcaaaagggcgtttgttagatattagataaataccgtgta	971
OY	964	AAGTCCTAGGACAGTGGCAGCGCAATATGGAGGACATTCATGAACATTTTTGGCATTAAT	1021
Db	972	aagtgccataggcagtgccagcgaacaatgagggagcatccaatgaacatctttttgcataataa	1031
OY	1024	CCAAAAAATAACTGTATTCATATAAAACCTGCATCCACATGAATATTC	1072
Db	1032	ccaaaaataactcgtatcaataaaacttgatccaaatbaatttc	1080

RESULT 10
AAE81788
ID AAE81788 standard; cDNA; 1315 BP.
XX
XX AAE81788;
AC
XX
XX 12-JUN-2001 (first entry)
DT
XX
DE Human secreted protein gene 2 SPO ID NO:12.

OS Homo sapiens.
PN WO200112775-A2.
PD 22-FEB-2001.
XX
XX 16-AUG-2000; 2000WO-US22325.
XX
XX 17-AUG-1999; 99US-0149182.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
PA
PI Rosen CA, Ni J, Florence KA, Fiscella M, Wei P, Baker KP;
PI Birse CE, Young PE, Komatsuolis GA, Moore PA, Soppet DR;
XX
XX WPI; 2001-147550/15.
DR P-PSDB; AAB74734.
XX
PT Nucleic acids encoding 25 human secreted polypeptides, useful for
PT preventing, diagnosing and/or treating e.g. cancers, Parkinson's
PT disease and diabetic retinopathy -
XX
XX
PS Claim 1; Page 441-442; 485pp; English.
XX
XX AABF187 to AAF81817 encode the human secreted proteins given in AAB74733
CC to AAB74772. Human secreted proteins can have activities based on the
CC tissues and cells they are expressed in. Example of activities include:
CC immunomodulatory; antisclerotic; dermatological; immunosuppressive;
CC antiinflammatory; anti-HIV; immunostimulant; cytostatic; cardiant;
CC vascular; anti-angiogenic; ophthalmological; neuroprotectant; nootropic;
CC antiviral; antialzheimers; antiparkinsonian; antimicrobial; and
CC vulnery. Human secreted proteins can be used in gene therapy and
CC vaccine. Human secreted protein nucleotide sequences (NAMI) and proteins
CC (PEP1) may be used in the prevention, diagnosis and treatment of diseases
CC associated with inappropriate polypeptide expression. For example, NAMI
CC and PEP1 may be used to treat disorders associated with decreased
CC expression by rectifying mutations or deletions in a patients genome

Db 384 TTGGCATATATAAACCAAAAATTAACCTGTATTCATATAAAAACTTGATCCATCAATGAATTT 325
Qy 1072 C 1072
Db 324 C 324

RESULT 13
AAH72087/c
ID AAH72087 standard; cDNA; 468 BP.
XX
AC AAH72087;
XX
DT 19-SEP-2001 (first entry)
XX
DE Human cervical cancer marker nucleic acid 3361.
XX
KM Cervical cancer; cytostatic; pre-malignant condition; gene therapy; ss.
XX
OS Homo sapiens.
XX
PN WC0200142467-A2.
XX
PD 14-JUN-2001.
XX
PF 08-DEC-2000; 2000MO-US33312.
XX
PR 08-DEC-1999; 99US-0169681.
PR 21-DEC-1999; 99US-0171350.
PR 14-MAR-2000; 2000US-0189315.
PR 12-MAY-2000; 2000US-0203791.
PR 09-JUN-2000; 2000US-0210600.
PR 21-JUL-2000; 2000US-0220114.
XX
PA (MILL-) MILENNIUM PREDICTIVE MEDICINE INC.
XX
PI Schlegel R, Deeds J, Berger A, Zhao X;
XX
DR WPI; 2001-375006/39.
XX
PT New isolated nucleic acid for diagnosing and treating cervical cancer
PT and for assessing and detecting compounds for treating the cancer -
XX
PS Claim 1; Page 652; 1051pp; English.
XX
CC The invention relates to novel genes (AAH68727-AAH73383) associated with
CC cervical cancer with cytostatic activity. The nucleic acids and encoded
CC polypeptides are useful: to assess if a patient is afflicted with
CC cervical cancer or has a pre-malignant condition; to monitor the
CC progression of cervical cancer or a premalignant condition in a patient;
CC and to select and/or assess the efficacy of a compound or therapy for
CC inhibiting cervical cancer in a patient. The nucleic acids may also be
CC useful for gene therapy.
XX
SQ Sequence 468 BP; 131 A; 84 C; 94 G; 159 T; 0 other;

Query Match 33.2%; Score 356; DB 22; Length 468;
Best Local Similarity 100.0%; Pred. No. 2.5e-167;
Matches 356; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 717 CCATCATCTTCTGGGGGAAAAATCTAGTATTTTGTATTTGGAATCTTACAGCAACAA 776
Db 468 CCATCATCTTCTGGGGGAAAAATCTAGTATTTTGTATTTGGAATCTTACAGCAACAA 409
Qy 777 ATGGAGCTCTGGCCCATGAGAGCTCTTGACAGTAATCACCACGCCGATACGAAGTC 836
Db 408 ATGGAGCTCTGGCCCATGAGAGCTCTTGACAGTAATCACCACGCCGATACGAAGTC 349
Qy 837 TTGCAACAACAAAATGTGTGGCAATAGAACTATATCAAGCAATATCTCCACCACCAAGC 896
Db 348 TTGCAACAACAAAATGTGTGGCAATAGAACTATATCAAGCAATATCTCCACCACCAAGC 269

Qy 897 TTCTGTAACCTGGGACCAATGATTAACCTCATAGGCGCTGTGTGAGAGTTAGATGAATA 956
Db 288 TTCTGTAACCTGGGACCAATGATTAACCTCATAGGCGCTGTGTGAGAGTTAGATGAATA 229
Qy 957 CCTGTGAAAGTGGCTAGGCAAGTGGCCAGCCAAATAGAGAGCATTCATGAACATTTTTCG 1016
Db 228 CCTGTGAAAGTGGCTAGGCAAGTGGCCAGCCAAATAGAGAGCATTCATGAACATTTTTCG 169
Qy 1017 ATATTAACCAAAAATTAACCTGTATTCATATAAAAACTTGATCCATCAATGAATTC 1072
Db 168 ATATTAACCAAAAATTAACCTGTATTCATATAAAAACTTGATCCATCAATGAATTC 113

RESULT 14
AAH1842/c
ID AAH1842 standard; cDNA; 528 BP.
XX
AC AAH1842;
XX
DT 26-JUN-2001 (first entry)
XX
DE Human cDNA clone (3'-primer) SEQ ID NO:8677.
XX
KM Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX
OS Homo sapiens.
XX
PN EP1074617-A2.
XX
PD 07-FEB-2001.
XX
PF 28-JUL-2000; 2000BP-0116126.
XX
PR 29-JUL-1999; 99JP-0248036.
PR 27-AUG-1999; 99JP-0300253.
PR 11-JAN-2000; 2000JP-0118776.
PR 02-MAY-2000; 2000JP-0183767.
PR 09-JUN-2000; 2000JP-0241899.
XX
PA (HELI-) HELIX RES INST.
XX
PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
DR WPI; 2001-318749/34.
XX
PT Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
XX full-length cDNAs -
XX
PS Claim 3; SEQ ID 8677; 2537pp + CD ROM; English.

CC The present invention describes primer sets for synthesizing 5602
CC full-length cDNAs defined in the specification. Where a primer set
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
CC to the complementary strand of a polynucleotide which comprises one of
CC the 5602 nucleotide sequences defined in the specification, where the
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC of an oligonucleotide comprising a sequence complementary to the
CC complementary strand of a polynucleotide which comprises a 5'-end
CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesizing polynucleotides,
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632

CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.
XX
SQ Sequence 528 BP; 153 A; 89 C; 108 G; 177 T; 1 other;

Query Match 32.0%; Score 343; DB 22; Length 528;
Best Local Similarity 100.0%; Pred. No. 7.8e-161;
Matches 343; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 730 GGGGAAAAATCTAGTATTGATTATTGATCTTACGACAAATAGCACTCTG 789
DB 485 GGGGAAAAATCTAGTATTGATTATTGATCTTACGACAAATAGCACTCTG 426
OY 790 GCCAATGAGAGCTCTGACCAATGATCCAGCCGATCGAAGCTTGGCCACAAAA 849
DB 425 GCCAATGAGAGCTCTGACCAATGATCCAGCCGATCGAAGCTTGGCCACAAAA 366
OY 850 TGTGTGGCAATAGATATATCAAGCAATAATCTCCACCCAGGCTTGTAAACTGG 909
DB 365 TGTGTGGCAATAGATATATCAAGCAATAATCTCCACCCAGGCTTGTAAACTGG 306
OY 910 GACCAATGATTTACCTCATAGGCGTGTGAGAGATTAGATGAATACCTGTGAAGTGC 969
DB 305 GACCAATGATTTACCTCATAGGCGTGTGAGAGATTAGATGAATACCTGTGAAGTGC 246
OY 970 CTAGCAGATGCGCAGCAATAGAGAGCATTCATGAACTTTTTCATATTAACCAAAA 1029
DB 245 CTAGCAGATGCGCAGCAATAGAGAGCATTCATGAACTTTTTCATATTAACCAAAA 186
OY 1030 AATACTTGTATTCATATTAATAAATCTGCATCAACATGAATTC 1072
DB 185 AATACTTGTATTCATATTAATAAATCTGCATCAACATGAATTC 143

RESULT 15

AAZ65182

AAZ65182;

05-APR-2000 (first entry)

Probe specific for human PRO828.

Membrane-bound polypeptide: PRO polypeptide; LDL receptor; TIE ligand;
pharmaceutical; receptor immunoadhesin; gene mapping; probe; ss.

Homo sapiens.

WO9963088-A2.

09-DEC-1999.

02-JUN-1999; 99WO-US12252.

02-JUN-1998; 98US-0087607.

02-JUN-1998; 98US-0087559.

03-JUN-1998; 98US-0087827.

04-JUN-1998; 98US-0088021.

04-JUN-1998; 98US-0088025.

04-JUN-1998; 98US-0088028.

04-JUN-1998; 98US-0088029.

04-JUN-1998; 98US-0088030.

04-JUN-1998; 98US-0088033.

04-JUN-1998; 98US-0088326.

05-JUN-1998; 98US-0088202.

05-JUN-1998; 98US-0088212.

05-JUN-1998; 98US-0088217.

09-JUN-1998; 98US-0088655.

10-JUN-1998; 98US-0088722.

PR 10-JUN-1998; 98US-0088730.
PR 10-JUN-1998; 98US-0088734.
PR 10-JUN-1998; 98US-0088738.
PR 10-JUN-1998; 98US-0088740.
PR 10-JUN-1998; 98US-0088741.
PR 10-JUN-1998; 98US-0088742.
PR 10-JUN-1998; 98US-0088810.
PR 10-JUN-1998; 98US-0088811.
PR 10-JUN-1998; 98US-0088824.
PR 10-JUN-1998; 98US-0088825.
PR 10-JUN-1998; 98US-0088826.
PR 11-JUN-1998; 98US-0088858.
PR 11-JUN-1998; 98US-0088861.
PR 11-JUN-1998; 98US-0088863.
PR 11-JUN-1998; 98US-0088876.
PR 12-JUN-1998; 98US-0088909.
PR 12-JUN-1998; 98US-0089105.
PR 16-JUN-1998; 98US-0089440.
PR 16-JUN-1998; 98US-0089512.
PR 16-JUN-1998; 98US-0089514.
PR 17-JUN-1998; 98US-0089532.
PR 17-JUN-1998; 98US-0089538.
PR 17-JUN-1998; 98US-0089598.
PR 17-JUN-1998; 98US-0089599.
PR 17-JUN-1998; 98US-0089600.
PR 17-JUN-1998; 98US-0089653.
PR 18-JUN-1998; 98US-0089801.
PR 18-JUN-1998; 98US-0089907.
PR 18-JUN-1998; 98US-0089908.
PR 19-JUN-1998; 98US-0089947.
PR 19-JUN-1998; 98US-0089948.
PR 19-JUN-1998; 98US-0089952.
PR 22-JUN-1998; 98US-0090246.
PR 22-JUN-1998; 98US-0090252.
PR 22-JUN-1998; 98US-0090254.
PR 23-JUN-1998; 98US-0090349.
PR 23-JUN-1998; 98US-0090355.
PR 24-JUN-1998; 98US-0090429.
PR 24-JUN-1998; 98US-0090431.
PR 24-JUN-1998; 98US-0090435.
PR 24-JUN-1998; 98US-0090444.
PR 24-JUN-1998; 98US-0090445.
PR 24-JUN-1998; 98US-0090461.
PR 24-JUN-1998; 98US-0090472.
PR 24-JUN-1998; 98US-0090535.
PR 24-JUN-1998; 98US-0090538.
PR 24-JUN-1998; 98US-0090540.
PR 24-JUN-1998; 98US-0090557.
PR 25-JUN-1998; 98US-0090676.
PR 25-JUN-1998; 98US-0090678.
PR 25-JUN-1998; 98US-0090688.
PR 25-JUN-1998; 98US-0090690.
PR 25-JUN-1998; 98US-0090691.
PR 25-JUN-1998; 98US-0090694.
PR 25-JUN-1998; 98US-0090695.
PR 25-JUN-1998; 98US-0090696.
PR 26-JUN-1998; 98US-0090862.
PR 26-JUN-1998; 98US-0090863.
PR 01-JUL-1998; 98US-0091358.
PR 01-JUL-1998; 98US-0091360.
PR 01-JUL-1998; 98US-0091544.
PR 02-JUL-1998; 98US-0091478.
PR 02-JUL-1998; 98US-0091486.
PR 02-JUL-1998; 98US-0091519.
PR 02-JUL-1998; 98US-0091626.
PR 02-JUL-1998; 98US-0091628.
PR 02-JUL-1998; 98US-0091633.
PR 02-JUL-1998; 98US-0091646.
PR 02-JUL-1998; 98US-0091673.
PR 02-JUL-1998; 98US-0091978.
PR 07-JUL-1998; 98US-0091982.
PR 09-JUL-1998; 98US-0092182.
PR 10-JUL-1998; 98US-0092472.

PR 20-JUL-1998; 98US-0093339.
 PR 30-JUL-1998; 98US-0094651.
 PR 04-AUG-1998; 98US-0095282.
 PR 04-AUG-1998; 98US-0095285.
 PR 04-AUG-1998; 98US-0095301.
 PR 04-AUG-1998; 98US-0095302.
 PR 04-AUG-1998; 98US-0095318.
 PR 04-AUG-1998; 98US-0095321.
 PR 04-AUG-1998; 98US-0095322.
 PR 10-AUG-1998; 98US-0095916.
 PR 10-AUG-1998; 98US-0095929.
 PR 10-AUG-1998; 98US-0096012.
 PR 11-AUG-1998; 98US-0096143.
 PR 11-AUG-1998; 98US-0096146.
 PR 12-AUG-1998; 98US-0096329.
 PR 17-AUG-1998; 98US-0096757.
 PR 17-AUG-1998; 98US-0096766.
 PR 17-AUG-1998; 98US-0096768.
 PR 17-AUG-1998; 98US-0096773.
 PR 17-AUG-1998; 98US-0096791.
 PR 17-AUG-1998; 98US-0096867.
 PR 17-AUG-1998; 98US-0096891.
 PR 17-AUG-1998; 98US-0096894.
 PR 17-AUG-1998; 98US-0096895.
 PR 17-AUG-1998; 98US-0096897.
 PR 18-AUG-1998; 98US-0096949.
 PR 18-AUG-1998; 98US-0096950.
 PR 18-AUG-1998; 98US-0096959.
 PR 18-AUG-1998; 98US-0096960.
 PR 18-AUG-1998; 98US-0097022.
 PR 19-AUG-1998; 98US-0097141.
 PR 20-AUG-1998; 98US-0097218.
 PR 24-AUG-1998; 98US-0097661.
 PR 26-AUG-1998; 98US-0097951.
 PR 26-AUG-1998; 98US-0097952.
 PR 26-AUG-1998; 98US-0097954.
 PR 26-AUG-1998; 98US-0097955.
 PR 26-AUG-1998; 98US-0097971.
 PR 26-AUG-1998; 98US-0097978.
 PR 26-AUG-1998; 98US-0097978.
 PR 26-AUG-1998; 98US-0097979.
 PR 26-AUG-1998; 98US-0097986.
 PR 26-AUG-1998; 98US-0098014.
 PR 31-AUG-1998; 98US-0098525.
 PR 16-SEP-1998; 98US-0100634.
 PR 12-JAN-1999; 99US-0115565.
 XX (GETH) GENENTECH INC.
 PA Baker K, Chen J, Goddard A, Gurney AL, Smith V, Watanabe CK;
 PI Wood WI, Yuan J;
 XX WPI: 2000-072883/06.
 DR
 XX
 .PT
 XX
 PS
 Example 50; Page 418: 822pp: English.
 Membrane-bound proteins and related nucleotide sequences -
 The invention provides membrane-bound PRO polypeptides and polynucleotides encoding them. The PRO sequences of the invention were identified based on extracellular domain homology screening. The PRO sequences have homology with proteins including LDL receptors, TIR ligands and various enzymes. The membrane-bound proteins and receptor molecules are useful as pharmaceutical and diagnostic agents. Receptor immunoadhesins, for instance, can be used as therapeutic agents to block receptor-ligand interactions. The membrane-bound proteins can also be employed for screening of potential peptide or small molecule inhibitors of the relevant receptor/ligand interaction. The PRO encoding sequences are useful as hybridization probes, in chromosome and gene mapping and in the generation of antisense RNA and DNA. PRO nucleic acid sequences will also be useful for the preparation of PRO polypeptides, especially by recombinant techniques.

SQ Sequence 50 BP; 11 A; 12 C; 18 G; 9 T; 0 other;
 Query Match 4.7%; Score 50; DB 21; Length 50;
 Best Local Similarity 100.0%; Pred. No. 1,1e-14;
 Matches 50; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 118 CAACATCCGGGCGCAACTGTGTCTGCTGAGAGTACCGCGGATCGGTCT 167
 DB 1 caacatccgggccaactgtgtctgctgagagatcaccgcgatcgggtct 50
 Search completed: August 25, 2002, 07:18:34
 Job time: 3889 sec

Mon Aug 26 08:01:42 2002

us-09-811-118-2.oli.rng

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